# New England Biolabs Certificate of Analysis 

Product Name:
Catalog Number:
Concentration:
Unit Definition:
Packaging Lot Number:
Expiration Date:
Storage Temperature:
Storage Conditions:
Specification Version:
dGTP Solution
N0442S
100 mM
N/A
10147948
12/2023
$-20^{\circ} \mathrm{C}$
Supplied in Ultrapure water as a sodium salt ( pH 7.5 )
PS-N0442S v2.0
dGTP Solution Component List

| NEB Part Number | Component Description | Lot Number | Individual QC Result |
| :--- | :--- | :--- | :---: |
| N0442SVIAL | dGTP | 10132189 | Pass |


| Assay Name/Specification | Lot \# 10147948 |
| :---: | :---: |
| qPCR DNA Contamination (E. coli Genomic) <br> A minimum of $1 \mu$ l of dGTP Solution is screened for the presence of E. coli genomic DNA using SYBR® Green qPCR with primers specific for the E. coli 16S rRNA locus. Results are quantified using a standard curve generated from purified E. coli genomic DNA. The measured level of E . coli genomic DNA contamination is $\leq 1 \mathrm{E}$. coli genome. | Pass |
| RNase Activity (Extended Digestion) <br> A $10 \mu$ reaction in NEBuffer 4 containing 40 ng of a 300 base single-stranded RNA and a minimum of $1 \mu$ of dGTP Solution is incubated at $37^{\circ} \mathrm{C}$. After incubation for 16 hours, $>90 \%$ of the substrate RNA remains intact as determined by gel electrophoresis using fluorescent detection. | Pass |
| Endonuclease Activity (Nicking) <br> A $50 \mu \mathrm{l}$ reaction in NEBuffer 2 containing $1 \mu \mathrm{~g}$ of supercoiled PhiX174 DNA and a minimum of $1 \mu$ l of dGTP Solution incubated for 4 hours at $37^{\circ} \mathrm{C}$ results in $<10 \%$ conversion to the nicked form as determined by agarose gel electrophoresis. | Pass |
| PCR Amplification ( 5.0 kb Lambda, dNTPs) <br> A $50 \mu \mathrm{l}$ reaction in ThermoPol® Reaction Buffer in the presence of $200 \mu \mathrm{M}$ dATP, dTTP, dCTP, and dGTP and $0.2 \mu \mathrm{M}$ primers containing 1 ng Lambda DNA with 1.25 units of Taq DNA Polymerase for 25 cycles of PCR amplification results in the expected 5.0 kb | Pass |


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| product. |  |
| Phosphatase Activity (pNPP) <br> A $200 \mu \mathrm{l}$ reaction in 1 M Diethanolamine, $\mathrm{pH} 9.8,0.5 \mathrm{mM} \mathrm{MgCl} 2$ containing 2.5 mM p-Nitrophenyl Phosphate (pNPP) and a minimum of $16 \mu$ of dGTP Solution incubated for 4 hours at $37^{\circ} \mathrm{C}$ yields $<0.0001$ unit of alkaline phosphatase activity as determined by spectrophotometric analysis. | Pass |
| PCR Amplification ( $\mathbf{2 . 0} \mathbf{~ k b}$ Lambda, dNTPs) <br> A $50 \mu \mathrm{l}$ reaction in ThermoPol® Reaction Buffer in the presence of $200 \mu \mathrm{M}$ dATP, dTTP, dCTP, and dGTP and $0.2 \mu \mathrm{M}$ primers containing 1 ng Lambda DNA with 1.25 units of Taq DNA Polymerase for 25 cycles of PCR amplification results in the expected 2.0 kb product. | Pass |
| PCR Amplification ( 0.5 kb Lambda, dNTPs) <br> A $50 \mu \mathrm{l}$ reaction in ThermoPol® Reaction Buffer in the presence of $200 \mu \mathrm{M}$ dATP, dTTP, dCTP, and dGTP and $0.2 \mu \mathrm{M}$ primers containing 1 ng Lambda DNA with 1.25 units of Taq DNA Polymerase for 25 cycles of PCR amplification results in the expected 0.5 kb product. | Pass |
| Non-Specific DNase Activity (16 Hour) <br> A $50 \mu$ l reaction in NEBuffer 2 containing $1 \mu \mathrm{~g}$ of T3 or T7 DNA in addition to a reaction containing Lambda-HindIII DNA and a minimum of $4 \mu$ I of dGTP Solution incubated for 16 hours at $37^{\circ} \mathrm{C}$ results in a DNA pattern free of detectable nuclease degradation as determined by agarose gel electrophoresis. | Pass |
| Physical Purity (HPLC) <br> dGTP Solution is $\geq 99 \%$ pure as determined by HPLC analysis. | Pass |

This product has been tested and shown to be in compliance with all specifications.
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## Chiastic Vazquez

Christie Vazquez
Production Scientist
05 Apr 2022


Michael Tonello
Packaging Quality Control Inspector 05 Apr 2022

